

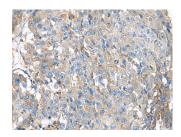
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

# **KIAA0556 Polyclonal Antibody**

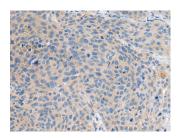
Catalog No.E-AB-52101ReactivityH,MStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsIHC,ELISAIsotypeIgG

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

# **Images**



Immunohistochemistry of paraffinembedded Human liver cancer tissue using KIAA0556 Polyclonal Antibody at dilution of 1:90(×200)



Immunohistochemistry of paraffinembedded Human cervical cancer tissue using KIAA0556 Polyclonal Antibody at dilution of 1:90(×200)

# **Immunogen Information**

**Immunogen** Synthetic peptide of human KIAA0556

**Gene Accession** NP056017 **Swissprot** O60303

**Synonyms** K0556,KIAA0556,Uncharacterized protein

KIAA0556

#### **Product Information**

**Buffer** PBS with 0.05% NaN3 and 40% Glycerol,pH7.4

**Purify** Antigen affinity purification

**Dilution** IHC 1:100-1:200, ELISA 1:5000-1:10000

# **Background**

This gene encodes a novel, evolutionarily conserved, ciliary protein. In human hTERT-RPE1 cells, the protein is found at the base of cilia, decorating the ciliary axoneme, and enriched at the ciliary tip. The protein binds to microtubules in vitro and regulates their stability when it is overexpressed. A null mutation in this gene has been associated with Joubert syndrome, a recessive disorder that is characterized by a distinctive mid-hindbrain and cerebellar malformation and is also often associated with wider ciliopathy symptoms. Consistently, in a serumstarvation ciliogenesis assay, human fibroblast cells derived from patients with the mutation display a reduced number of ciliated cells with abnormally long cilia.